

ALLERGY RELIEF- diphenhydramine hcl liquid
LLORENS PHARMACEUTICALS INTERNATIONAL DIVISION

Llorens-AllergyChild 837

Active ingredient (in each 5 mL teaspoonful)

Diphenhydramine HCl 12.5 mg

Purpose

Antihistamine

Uses

- Temporarily relieves these symptoms of hay fever or other upper respiratory allergies:
- runny nose
- sneezing
- itching of the nose or throat
- itchy, watery eyes

Warnings

Do not use

- with any other product containing diphenhydramine, including one used on the skin
- to make a child sleepy

Ask a doctor before use if you have

- a breathing problem such as chronic bronchitis
- glaucoma

Ask a doctor or pharmacist before use if the child is

taking sedatives or tranquilizers.

When using this product

- **do not exceed recommended dose**
- excitability may occur, especially in children
- marked drowsiness may occur
- sedatives, and tranquilizers may increase drowsiness

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

- take every 4 to 6 hours as needed, or as directed by a doctor
- do not take more than 6 doses in 24 hours
- use dosing cup provided.

Age	Dosage
Children 6 to 11 years	5 mL to 10 mL

Children 2 to 5 years
Children under 2 years

Do not use unless directed by a doctor
Do not use

Inactive ingredients: Artificial and natural cherry flavor, citric acid, FD&C #40, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate, sucralose and sucrose.

Questions or comments? 1-866-595-5598

Drug Facts

Active Ingredient (in each 5 mL teaspoonful) Purpose
Diphenhydramine HCl 12.5 mg Antihistamine

Uses
Temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: • runny nose • sneezing • itching of the nose or throat • itchy, watery eyes

WARNINGS
Do not use • with any other product containing diphenhydramine, including one used on the skin • to make a child sleepy
Ask a doctor before use if the child has • a breathing problem such as chronic bronchitis • glaucoma
Ask a doctor or pharmacist before use if the child is taking sedatives or tranquilizers.

When using this product

- do not exceed recommended dose
- excitability may occur, especially in children
- marked drowsiness may occur
- sedatives and tranquilizers may increase drowsiness

*This product is not manufactured or distributed by the owner of the registered trademark Benadryl®.

Llorens Care

NDC: 54859-837-04

NON-HABIT FORMING

Compare to Children's Benadryl® Active Ingredients*

children's **allergy relief**
Diphenhydramine HCl 12.5 mg per 5 mL

Relief of:

- Runny Nose
- Sneezing
- Itchy, Watery Eyes
- Itchy Throat or Nose

Dye Free & Alcohol Free

Cherry Flavor

4 FL OZ (118 mL)

Drug Facts (continued)

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

- take every 4 to 6 hours, or as directed by a doctor.
- do not take more than 6 doses in 24 hours.
- use dosing cup provided.

Age	Dosage
Children 6 to 11 years	5 mL to 10 mL
Children 2 to 5 years	Do not use unless directed by a doctor
Children under 2 years	Do not use

Other information
TAMPER EVIDENT: Do not use if foil seal over bottle opening is torn, broken, or missing. • store at room temperature 15°-30°C (59°-86°F) • do not freeze • protect from light

Inactive ingredients: Artificial and natural cherry flavor, citric acid, FD&C Red #40, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate, sucralose, and sucrose.

Questions and comments? 1-866-595-5598

Manufactured by:
Llorens Pharmaceutical International Division, Inc. CODE #: L-191
Miami, FL 33147 REV.: 04/23
www.LlorensPharm.com

LOT #: _____
EXP DATE: _____

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ALLERGY RELIEF

diphenhydramine hcl liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54859-837
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII: 8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	12.5 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

SUCROSE (UNII: C151H8M554)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54859-837-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	12/01/2024	

Labeler - LLORENS PHARMACEUTICALS INTERNATIONAL DIVISION (037342305)

Revised: 12/2025

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